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NOTICE OF ALLOWANCE AND FEE(S) DUE

23347 7590 03/09/2009

GLAXOSMITHKLINE
CORPORATE INTELLECTUAL PROPERTY, MAI B482
FIVE MOORE DR., PO BOX 13398
RESEARCH TRIANGLE PARK, NC 27709-3398

EXAMINER	
RAO, DEEPAK R	
ART UNIT	PAPER NUMBER
1624	

DATE MAILED: 03/09/2009

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/560,502	12/13/2005	David Harold Drewry	PR60317USW	4610

TITLE OF INVENTION: CHEMICAL COMPOUNDS

APPLN. TYPE	SMALL ENTITY	ISSUE FEE DUE	PUBLICATION FEE DUE	PREV. PAID ISSUE FEE	TOTAL FEE(S) DUE	DATE DUE
nonprovisional	NO	$1510	$300	$0	$1810	06/09/2009

THE APPLICATION IDENTIFIED ABOVE HAS BEEN EXAMINED AND IS ALLOWED FOR ISSUANCE AS A PATENT. PROSECUTION ON THE MERITS IS CLOSED. THIS NOTICE OF ALLOWANCE IS NOT A GRANT OF PATENT RIGHTS. THIS APPLICATION IS SUBJECT TO WITHDRAWAL FROM ISSUE AT THE INITIATIVE OF THE OFFICE OR UPON PETITION BY THE APPLICANT. SEE 37 CFR 1.313 AND MPEP 1308.

THE ISSUE FEE AND PUBLICATION FEE (IF REQUIRED) MUST BE PAID WITHIN THREE MONTHS FROM THE MAILING DATE OF THIS NOTICE OR THIS APPLICATION SHALL BE REGARDED AS ABANDONED. THIS STATUTORY PERIOD CANNOT BE EXTENDED. SEE 35 U.S.C. 151. THE ISSUE FEE DUE INDICATED ABOVE DOES NOT REFLECT A CREDIT FOR ANY PREVIOUSLY PAID ISSUE FEE IN THIS APPLICATION. IF AN ISSUE FEE HAS PREVIOUSLY BEEN PAID IN THIS APPLICATION (AS SHOWN ABOVE), THE RETURN OF PART B OF THIS FORM WILL BE CONSIDERED A REQUEST TO REAPPLY THE PREVIOUSLY PAID ISSUE FEE TOWARD THE ISSUE FEE NOW DUE.

HOW TO REPLY TO THIS NOTICE:

I. Review the SMALL ENTITY status shown above.

If the SMALL ENTITY is shown as YES, verify your current SMALL ENTITY status:

A. If the status is the same, pay the TOTAL FEE(S) DUE shown above.

B. If the status above is to be removed, check box 5b on Part B - Fee(s) Transmittal and pay the PUBLICATION FEE (if required) and twice the amount of the ISSUE FEE shown above, or

If the SMALL ENTITY is shown as NO:

A. Pay TOTAL FEE(S) DUE shown above, or

B. If applicant claimed SMALL ENTITY status before, or is now claiming SMALL ENTITY status, check box 5a on Part B - Fee(s) Transmittal and pay the PUBLICATION FEE (if required) and 1/2 the ISSUE FEE shown above.

II. PART B - FEE(S) TRANSMITTAL, or its equivalent, must be completed and returned to the United States Patent and Trademark Office (USPTO) with your ISSUE FEE and PUBLICATION FEE (if required). If you are charging the fee(s) to your deposit account, section "4b" of Part B - Fee(s) Transmittal should be completed and an extra copy of the form should be submitted. If an equivalent of Part B is filed, a request to reapply a previously paid issue fee must be clearly made, and delays in processing may occur due to the difficulty in recognizing the paper as an equivalent of Part B.

III. All communications regarding this application must give the application number. Please direct all communications prior to issuance to Mail Stop ISSUE FEE unless advised to the contrary.

IMPORTANT REMINDER: Utility patents issuing on applications filed on or after Dec. 12, 1980 may require payment of maintenance fees. It is patentee's responsibility to ensure timely payment of maintenance fees when due.

PTOL-85 (Rev. 08/07) Approved for use through 08/31/2010.

PART B - FEE(S) TRANSMITTAL

Complete and send this form, together with applicable fee(s), to: Mail **Mail Stop ISSUE FEE**
Commissioner for Patents
P.O. Box 1450
Alexandria, Virginia 22313-1450
or Fax (571)-273-2885

INSTRUCTIONS: This form should be used for transmitting the ISSUE FEE and PUBLICATION FEE (if required). Blocks 1 through 5 should be completed where appropriate. All further correspondence including the Patent, advance orders and notification of maintenance fees will be mailed to the current correspondence address as indicated unless corrected below or directed otherwise in Block 1, by (a) specifying a new correspondence address; and/or (b) indicating a separate "FEE ADDRESS" for maintenance fee notifications.

CURRENT CORRESPONDENCE ADDRESS (Note: Use Block 1 for any change of address)

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RESEARCH TRIANGLE PARK, NC 27709-3398

Note: A certificate of mailing can only be used for domestic mailings of the Fee(s) Transmittal. This certificate cannot be used for any other accompanying papers. Each additional paper, such as an assignment or formal drawing, must have its own certificate of mailing or transmission.

Certificate of Mailing or Transmission

I hereby certify that this Fee(s) Transmittal is being deposited with the United States Postal Service with sufficient postage for first class mail in an envelope addressed to the Mail Stop ISSUE FEE address above, or being facsimile transmitted to the USPTO (571) 273-2885, on the date indicated below.

(Depositor's name)
(Signature)
(Date)

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/560,502	12/13/2005	David Harold Drewry	PR60317USW	4610

TITLE OF INVENTION: CHEMICAL COMPOUNDS

APPLN. TYPE	SMALL ENTITY	ISSUE FEE DUE	PUBLICATION FEE DUE	PREV. PAID ISSUE FEE	TOTAL FEE(S) DUE	DATE DUE
nonprovisional	NO	$1510	$300	$0	$1810	06/09/2009

EXAMINER	ART UNIT	CLASS-SUBCLASS
RAO, DEEPAK R	1624	514-274000

1. Change of correspondence address or indication of "Fee Address" (37 CFR 1.363).

❑ Change of correspondence address (or Change of Correspondence Address form PTO/SB/122) attached.

❑ "Fee Address" indication (or "Fee Address" Indication form PTO/SB/47; Rev 03-02 or more recent) attached. **Use of a Customer Number is required.**

2. For printing on the patent front page, list

(1) the names of up to 3 registered patent attorneys or agents OR, alternatively,

(2) the name of a single firm (having as a member a registered attorney or agent) and the names of up to 2 registered patent attorneys or agents. If no name is listed, no name will be printed.

1 _____

2 _____

3 _____

3. ASSIGNEE NAME AND RESIDENCE DATA TO BE PRINTED ON THE PATENT (print or type)

PLEASE NOTE: Unless an assignee is identified below, no assignee data will appear on the patent. If an assignee is identified below, the document has been filed for recordation as set forth in 37 CFR 3.11. Completion of this form is NOT a substitute for filing an assignment.

(A) NAME OF ASSIGNEE (B) RESIDENCE: (CITY and STATE OR COUNTRY)

Please check the appropriate assignee category or categories (will not be printed on the patent) : ❑ Individual ❑ Corporation or other private group entity ❑ Government

4a. The following fee(s) are submitted:

❑ Issue Fee

❑ Publication Fee (No small entity discount permitted)

❑ Advance Order - # of Copies _____

4b. Payment of Fee(s): **(Please first reapply any previously paid issue fee shown above)**

❑ A check is enclosed.

❑ Payment by credit card. Form PTO-2038 is attached.

❑ The Director is hereby authorized to charge the required fee(s), any deficiency, or credit any overpayment, to Deposit Account Number _____ (enclose an extra copy of this form).

5. **Change in Entity Status** (from status indicated above)

❑ a. Applicant claims SMALL ENTITY status. See 37 CFR 1.27.

❑ b. Applicant is no longer claiming SMALL ENTITY status. See 37 CFR 1.27(g)(2).

NOTE: The Issue Fee and Publication Fee (if required) will not be accepted from anyone other than the applicant; a registered attorney or agent; or the assignee or other party in interest as shown by the records of the United States Patent and Trademark Office.

Authorized Signature _____ Date _____

Typed or printed name _____ Registration No. _____

This collection of information is required by 37 CFR 1.311. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, Virginia 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, Virginia 22313-1450.

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number.

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EXAMINER	
RAO, DEEPAK R	
ART UNIT	PAPER NUMBER
1624	

DATE MAILED: 03/09/2009

Determination of Patent Term Adjustment under 35 U.S.C. 154 (b)

(application filed on or after May 29, 2000)

The Patent Term Adjustment to date is 597 day(s). If the issue fee is paid on the date that is three months after the mailing date of this notice and the patent issues on the Tuesday before the date that is 28 weeks (six and a half months) after the mailing date of this notice, the Patent Term Adjustment will be 597 day(s).

If a Continued Prosecution Application (CPA) was filed in the above-identified application, the filing date that determines Patent Term Adjustment is the filing date of the most recent CPA.

Applicant will be able to obtain more detailed information by accessing the Patent Application Information Retrieval (PAIR) WEB site (http://pair.uspto.gov).

Any questions regarding the Patent Term Extension or Adjustment determination should be directed to the Office of Patent Legal Administration at (571)-272-7702. Questions relating to issue and publication fee payments should be directed to the Customer Service Center of the Office of Patent Publication at 1-(888)-786-0101 or (571)-272-4200.

Notice of Allowability	Application No.	Applicant(s)	
	10/560,502	DREWRY ET AL.	
	Examiner	Art Unit	
	Deepak Rao	1624	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address--

All claims being allowable, PROSECUTION ON THE MERITS IS (OR REMAINS) CLOSED in this application. If not included herewith (or previously mailed), a Notice of Allowance (PTOL-85) or other appropriate communication will be mailed in due course. **THIS NOTICE OF ALLOWABILITY IS NOT A GRANT OF PATENT RIGHTS.** This application is subject to withdrawal from issue at the initiative of the Office or upon petition by the applicant. See 37 CFR 1.313 and MPEP 1308.

1. ☒ This communication is responsive to *the amendment filed on December 24, 2008*.

2. ☒ The allowed claim(s) is/are *63-67, 73-74*.

3. ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
   a) ☐ All b) ☐ Some* c) ☐ None of the:
      1. ☐ Certified copies of the priority documents have been received.
      2. ☐ Certified copies of the priority documents have been received in Application No. _____ .
      3. ☐ Copies of the certified copies of the priority documents have been received in this national stage application from the International Bureau (PCT Rule 17.2(a)).

   * Certified copies not received: _____.

Applicant has THREE MONTHS FROM THE "MAILING DATE" of this communication to file a reply complying with the requirements noted below. Failure to timely comply will result in ABANDONMENT of this application.
THIS THREE-MONTH PERIOD IS NOT EXTENDABLE.

4. ☐ A SUBSTITUTE OATH OR DECLARATION must be submitted. Note the attached EXAMINER'S AMENDMENT or NOTICE OF INFORMAL PATENT APPLICATION (PTO-152) which gives reason(s) why the oath or declaration is deficient.

5. ☐ CORRECTED DRAWINGS (as "replacement sheets") must be submitted.
   (a) ☐ including changes required by the Notice of Draftsperson's Patent Drawing Review (PTO-948) attached
      1) ☐ hereto or 2) ☐ to Paper No./Mail Date _____.
   (b) ☐ including changes required by the attached Examiner's Amendment / Comment or in the Office action of Paper No./Mail Date _____.

   Identifying indicia such as the application number (see 37 CFR 1.84(c)) should be written on the drawings in the front (not the back) of each sheet. Replacement sheet(s) should be labeled as such in the header according to 37 CFR 1.121(d).

6. ☐ DEPOSIT OF and/or INFORMATION about the deposit of BIOLOGICAL MATERIAL must be submitted. Note the attached Examiner's comment regarding REQUIREMENT FOR THE DEPOSIT OF BIOLOGICAL MATERIAL.

Attachment(s)

1. ☐ Notice of References Cited (PTO-892)
2. ☐ Notice of Draftperson's Patent Drawing Review (PTO-948)
3. ☒ Information Disclosure Statements (PTO/SB/08), Paper No./Mail Date 20081030
4. ☐ Examiner's Comment Regarding Requirement for Deposit of Biological Material
5. ☐ Notice of Informal Patent Application
6. ☐ Interview Summary (PTO-413), Paper No./Mail Date _____ .
7. ☒ Examiner's Amendment/Comment
8. ☐ Examiner's Statement of Reasons for Allowance
9. ☐ Other _____.

/Deepak Rao/
Primary Examiner
Art Unit 1624

EXAMINER'S AMENDMENT

An examiner's amendment to the record appears below. Should the changes and/or additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be submitted no later than the payment of the issue fee.

Authorization for this examiner's amendment was given in a telephone interview with Ms. Kathryn Coulter on February 24, 2009.

The application has been amended as follows:

In the Claims:

In claim 63, last line, delete: ", solvate, or physiologically functional derivative".

In claim 73, line 5, delete "or double".

(Copy of claims 63 and 73 as amended is enclosed in Appendix)

Receipt is acknowledged of the Information Disclosure Statement filed on October 30, 2008 and a copy is enclosed herewith.

Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Deepak Rao whose telephone number is (571) 272-0672. The examiner can normally be reached on Monday-Friday from 8:00am to 5:00pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, James O. Wilson, can be reached at (571) 272-0661. The fax phone number for the organization where this application or proceeding is assigned is (571) 273-8300.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (571) 272-1600.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

/Deepak Rao/
Primary Examiner
Art Unit 1624

March 12, 2009

APPENDIX

Copy of Claims 63 and 73 as amended:

63. (Currently amended) A compound selected from the group consisting of:

4-(4-fluorophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-[3,4-bis(ethyloxy)phenyl]-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-4-[4-(methylsulfonyl)phenyl]-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-2-oxo-4-(3-thienyl)-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-4,6-dimethyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-4-(1-naphthalenyl)-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-4-(2-naphthalenyl)-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-{5-[(1*H*-indazol-5-ylamino)carbonyl]-6-methyl-2-oxo-1,2,3,4-tetrahydro-4-pyrimidinyl}benzoic acid;

4-(2,4-difluorophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-4-[3-(methyloxy)phenyl]-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-4-[2-(methyloxy)phenyl]-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-cyanophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

3-{5-[(1*H*-indazol-5-ylamino)carbonyl]-6-methyl-2-oxo-1,2,3,4-tetrahydro-4-pyrimidinyl}benzoic acid;

4-(2-fluorophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(3-chloro-4-fluorophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-{3-[(2-hydroxyethyl)oxy]phenyl}-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-bromo-2-thienyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-hydroxyphenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-chloro-2-fluorophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-4-{3-[(methylsulfonyl)amino]phenyl}-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-2-oxo-4-(6-quinoxalinyl)-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-[4-(aminosulfonyl)phenyl]-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-2-oxo-4-(2-quinolinyl)-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-[3-fluoro-4-(methyloxy)phenyl]-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(3-cyanophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(1*H*-imidazol-1-yl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-2-oxo-4-(3-quinolinyl)-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-2-oxo-4-[(*E*)-2-phenylethenyl]-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-2-oxo-4-[4-(trifluoromethyl)phenyl]-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-chlorophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-[4-(acetylamino)phenyl]-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(2-chlorophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(2,3-dihydro-1,4-benzodioxin-6-yl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(3-hydroxyphenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(8-hydroxy-2-quinolinyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-[3,4-bis(methyloxy)phenyl]-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-[2-(4-chlorophenyl)ethyl]-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-[3-(1*H*-imidazol-1-yl)phenyl]-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(3-chlorophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-[4-(aminocarbonyl)phenyl]-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-fluorophenyl)-*N*-1*H*-indazol-5-yl-6-(1-methylethyl)-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-fluorophenyl)-6-(2-furanyl)-*N*-1*H*-indazol-5-yl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-fluorophenyl)-*N*-1*H*-indazol-5-yl-1,6-dimethyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-1,6-dimethyl-4-(2-naphthalenyl)-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-4-(2-naphthalenyl)-2-thioxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-fluorophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-thioxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

N-1*H*-indazol-5-yl-6-methyl-4-(3-thienyl)-2-thioxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-fluorophenyl)-*N*-1*H*-indazol-5-yl-1,3,6-trimethyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-fluorophenyl)-*N*-1*H*-indazol-5-yl-6-methyl-2-oxo-1-(phenylmethyl)-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

4-(4-fluorophenyl)-*N*-1*H*-indazol-5-yl-*N*,6-dimethyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

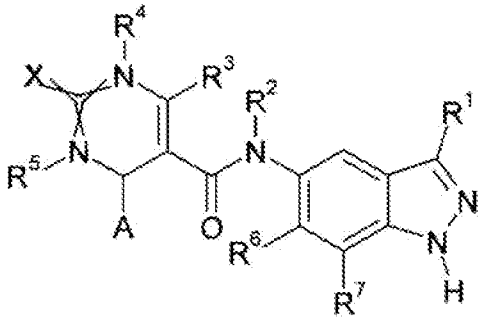
4-(4-fluorophenyl)-N-1H-indazol-5-yl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

1-ethyl-4-(4-fluorophenyl)-N-1H-indazol-5-yl-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide; and

N-(3-amino-1H-indazol-5-yl)-4-(4-fluorophenyl)-6-methyl-2-oxo-1,2,3,4-tetrahydro-5-pyrimidinecarboxamide;

or a salt~~, solvate, or physiologically functional derivative~~ thereof.

73. (Currently amended) A compound of Formula (I):



(I)

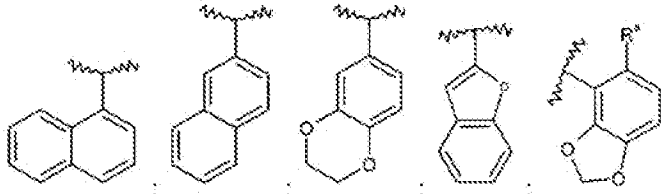
or a salt thereof:

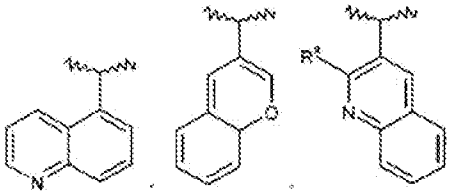
wherein:

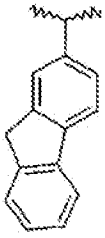
~~—~~ indicates a single ~~or double~~ bond;

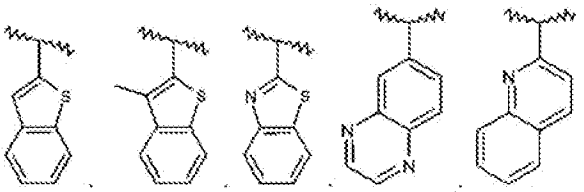
X is =O or =S;

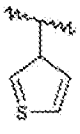
A is selected from $-CH_2OCH_2R'''$ where R''' is phenyl; cyclohexenylene;

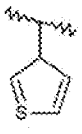

; ; ; ; where R^a is –H or $-OCH_3$; where R^a is –Cl;

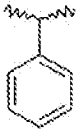

; ; where R^a is –H or –Cl;

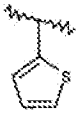

;

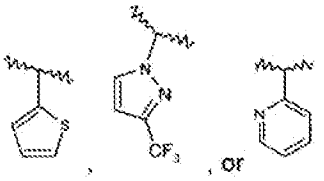

; ; ; ; optionally substituted with –OH;

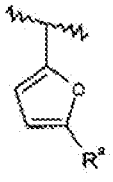

optionally substituted with –Br;

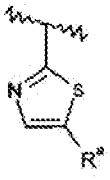

; or

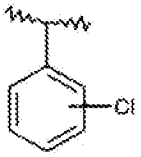

optionally substituted with one or more groups selected from –Cl, $-CF_3$, and methoxy;


optionally substituted with $-CH_3$, phenyl, –Cl, ethynyl substituted with phenyl,


, , or ;

 where R^a is $-CH_3$ or phenyl substituted with –Cl;

 where R^a is phenyl or $-SCH_3$;

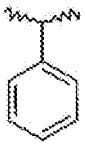
C_1-C_6 alkyl optionally substituted with aryl or  ;

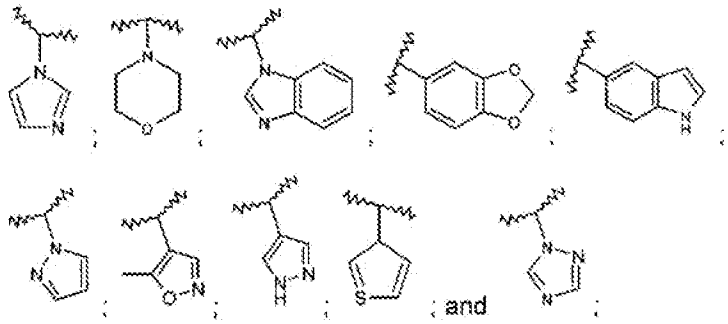
C_2-C_6 alkenyl optionally substituted with phenyl, where the phenyl is optionally substituted with $-NO_2$ or methoxy;

C_2-C_6 alkenyl optionally independently di-substituted with phenyl and/or –Cl;

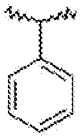
C_1-C_6 alkenyl optionally substituted with furanyl;

C_2-C_6 alkynyl optionally substituted with phenyl;

or A is  optionally substituted with one or more groups selected from –F; -Cl; –Br; -F; methoxy, ethoxy; $-S(O)_2CH_3$; $-N(H)S(O)_2CH_3$; $-S(O)_2NH_2$; $-C(O)NH_2$; -C(O)OH; -CN; -OH; $-O(CH_2)_rOH$, where r is 1, 2, 3, or 4; $-N(H)C(O)CH_3$; $-CF_3$, $-NO_2$; phenoxy, benzyloxy; $-OCF_3$; $-NR^aR^a$ where R^a is independently –H, $-CH_3$ or $-CH_2CH_3$;



; and



or A is substituted with phenyl wherein said phenyl is optionally substituted with one or more groups selected from–Cl, -F; $–CF_3$; methoxy or $-NR^aR^a$ where R^a is independently –H, $–CH_3$ or $–CH_2CH_3$; or $–C(O)CH_3$;

R^1 is –H, –NH(R'), –Cl, –Br, $–CH_3$, or phenyl optionally substituted with –F;

R' is –H or phenyl optionally substituted with one or more –F;

R^2 is –H or $–CH_3$ and R^3 is –H, C_1-C_3 alkyl, $–CH_2OCH_3$, or furanyl; or R^2 and R^3 together with the ring and atoms to which they are attached form a fused ring system;

R^4 is –H, $–CH_3$, $–CH_2CH_3$, or benzyl;

R^5 is –H or $–CH_3$;

R^6 is –H, –Cl, or –F; and

R^7 is –H or –Cl.